Claims:

1. A method for preparing a therapeutic aqueous disodium pamidronate solution comprising:

(a) preparing a slurry of pamidronic acid in water, and

(b) combining aqueous sodium hydroxide with said slurry in an about 2:1 molar ratio of sodium hydroxide to pamidronic acid; to yield a solution of disodium pamidronate having visual clarity and a pH of about 6.5;

(c) packaging said solution in a plurality of sealed containers to yield a plurality of liquid unit dosage forms of pamidronate.

2. The method of claim 1 wherein the slurry includes an effective stablizing amount of mannitol.

3. A unit dosage form comprising a disodium pamidronte solution prepared by the method of claim 1.

4. A unit dosage form comprising a disodium pamidronate solution prepared by the method of claim 2.

5. A vial or ampule comprising a unit dosage form of the solution of claim 3.

6. A vial or ampule comprising a unit dosage form of the solution of claim 4.

7. The vial or ampule of claim 5 wherein the solution is packaged under an inert atmosphere.

8. The dosage form of claim 3 wherein the container is free of CA^{+2} that can be sequestered by the disodium pamidronate.

9. The dosage form of claim 8 wherein the container is a plastic vial or ampule.

10. The dosage form of claim 8 wherein the container is a glass vial or ampule.